

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 275 651 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
15.01.2003 Bulletin 2003/03

(51) Int Cl.7: **C07D 495/14, C07D 519/00,
C09K 19/34, H01B 1/12**

(21) Application number: **02013425.0**

(22) Date of filing: **13.06.2002**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **09.07.2001 EP 01115742**

(71) Applicant: **MERCK PATENT GmbH
64293 Darmstadt (DE)**

(72) Inventors:
• **Farrand, Louise**
Blandford Forum, Dorset DT11 9ED (GB)

- **Thompson, Marcus**
Fordingbridge, Hampshire SP6 1RR (GB)
- **Giles, Mark**
Southampton SO15 2LE (GB)
- **Heeney, Martin**
Southampton SO14 6TQ (GB)
- **Tierney, Steven**
Southampton SO15 7QW (GB)
- **Shkunov, Maxim**
Southampton SO16 6SX (GB)
- **Sparrowe, David**
Bournemouth, Dorset BH6 5EJ (GB)
- **McCulloch, Iain**
Kings Somborne, Hampshire SO20 6PE (GB)

(54) **Thienothiophene derivatives**

(57) The invention relates to new thienothiophene derivatives, their use as semiconductors or charge transport materials, in optical, electrooptical or electronic devices like for example organic field effect transistors (FET or OFET) for thin film transistor liquid crystal displays and integrated circuit devices such as RFID tags,

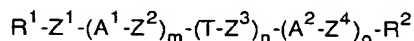
electroluminescent devices in flat panel displays, and in photovoltaic and sensor devices, and to a field effect transistor, light emitting device or ID tag comprising the thienothiophene derivatives.

EP 1 275 651 A2

with the ability to induce liquid crystal phase behaviour. The compounds or materials comprising mesogenic groups do not necessarily have to exhibit a liquid crystal phase themselves. It is also possible that they show liquid crystal phase behaviour only in mixtures with other compounds.

5 Summary of the Invention

[0013] One object of the invention are thienothiophene derivatives of formula I



wherein

15 R^1 and R^2 are independently of each other H, halogen, CN, NO_2 , straight chain, branched or cyclic alkyl with 1 to 20 C-atoms, which may be unsubstituted, mono- or poly-substituted by F, Cl, Br, I or CN, it being also possible for one or more non-adjacent CH_2 groups to be replaced, in each case independently from one another, by -O-, -S-, -NH-, - NR^0 -, - SiR^0R^{00} -, -CO-, -COO-, -OCO-, -OCO-O-, -S-CO-, -CO-S-, -CH=CH- or -C≡C- in such a manner that O and/or S atoms are not linked directly to one another,

20 R^0 and R^{00} are independently of each other H or alkyl with 1 to 12 C-atoms,

A^1 and A^2 are independently of each other an alicyclic or aromatic group that may also comprise one or more hetero atoms and one or more fused rings, and A^1 may also denote T,

25 Z^1 to Z^4 are independently of each other -O-, -S-, -CO-, -COO-, -OCO-, -O-COO-, -CO- NR^0 -, - NR^0 -CO-, -OCH₂-, -CH₂O-, -SCH₂-, -CH₂S-, -CF₂O-, -OCF₂-, -CF₂S-, -SCF₂-, -CH₂CH₂-, -CF₂CH₂-, -CH₂CF₂-, -CF₂CF₂-, -CH=N-, -N=CH-, -N=N-, -CH=CR⁰-, -CX¹=CX²-, -C≡C-, -CH=CH-COO-, -OCO-CH=CH- or a single bond,

30 X^1 and X^2 are independently of each other H, F, Cl or CN,

T is a group consisting of 3, 4, 5 or 6 fused thiophene rings which may also be mono- or polysubstituted by R^1 ,

35 m and o are independently of each other 0, 1, 2 or 3, and

n is 1, 2 or 3.

40 [0014] Another object of the invention is the use of compounds of formula I as semiconductors or charge transport materials, in particular in optical, electrooptical or electronic devices, like for example components of integrated circuitry, field effect transistors (FET) for example as thin film transistors in flat panel display applications or for Radio Frequency Identification (RFID) tags, or in semiconducting components for organic light emitting diode (OLED) applications such as electroluminescent displays or backlights of flat panel displays, for photovoltaic or sensor devices, as electrode materials in batteries, as photoconductors and for electrophotographic applications.

45 [0015] Another object of the invention is a field effect transistor, for example as a component of integrated circuitry, as a thin film transistor in flat panel display applications, or in an RFID tag, comprising one or more compounds of formula I.

50 [0016] Another object of the invention is a semiconducting component, for example in OLED applications like electroluminescent displays or backlights of flat panel displays, in photovoltaic or sensor devices, as electrode materials in batteries, as photoconductors and for electrophotographic applications, comprising one or more compounds of formula I.

[0017] Another object of the invention is a security marking or device comprising an RFID or ID tag or a FET according to the invention.

55 Detailed Description of the Invention

[0018] The compounds of formula I provide several advantages over prior art materials

- by adding substituent chains and other groups they can be made more soluble, thus being suitable for spin coating or solution coating techniques, rather than vacuum deposition, to prepare thin films for use e.g. in electronic devices such as transistors,
- 5 - they can be made mesogenic or liquid crystalline, thus exhibiting a higher degree of order that leads to particularly high charge carrier mobility, in particular when being aligned in their mesophase into macroscopically uniform orientation,
- 10 - they combine the properties of a semiconducting material with those of a mesogenic material to give novel materials with a rigid, planar conjugated core and a flexible chain to increase solubility and to decrease the melting point, which show high charge carrier mobility when being aligned in their mesophase.

[0019] The compounds of formula I are useful as charge transport semiconductors in that they have high carrier mobilities. In particular, the introduction of alkyl side chains to the thienothiophene core improves the solubility and therefore the solution processibility of the compounds of formula I.

[0020] Particularly preferred are mesogenic or liquid crystalline compounds of formula I, wherein T is a mesogenic group. These compounds are particularly useful as semiconductors or charge transport materials, as they can be processed while in the highly ordered mesophase morphology, and readily aligned by conventional techniques in a preferred direction. Both smectic and nematic mesophase ordering allows close packing of molecular pi-electron systems, which maximises intermolecular charge transfer which occurs through a hopping mechanism between adjacent molecules. In this way charge trap sites at grain boundaries between regions of different orientation are reduced.

[0021] Thus, another object of the invention is a liquid crystal mixture comprising one or more compounds of formula I and optionally comprising one or more further compounds, wherein at least one of the compounds of formula I and the further compounds is mesogenic or liquid crystalline.

[0022] It is also possible to mix compounds of formula I with other mesogenic or liquid crystal monomers that are known from prior art, in order to induce or enhance liquid crystal phase behaviour.

[0023] It is also possible to mix compounds of formula I with liquid crystal polymers or with polymerisable compounds which have one or more polymerisable groups and which may also be mesogenic or liquid crystalline. For example, if the compounds of formula I are mixed with one or more polymerisable mesogenic or liquid crystal compounds, the oriented as described above can be permanently "frozen-in" by polymerising the mesogens, which can also create a structure with long range order, or "monodomain". Formation of a monodomain also maximises charge transfer by eliminating charge trap sites at grain boundaries, while the polymerisation also improves the mechanical properties of the film. Further, by crosslinking the mesogens, a highly stable structure results, which has an additional advantage of being impervious to subsequent processing solvents during device fabrication, thus allowing a wider range of solvents to be used in deposition of the next layer of the device by solution techniques. In addition, it is often observed that this crosslinking further densifies the film, leading to smaller intermolecular distances and improved charge transport.

[0024] Thus, another object of the invention is a polymerisable liquid crystal mixture comprising one or more compounds of formula I and comprising one or more further polymerisable compounds, wherein the further polymerisable compounds may also be mesogenic or liquid crystalline.

[0025] Polymerisable mesogenic compounds that are suitable as comonomers are known in prior art and disclosed for example in WO 93/22397; EP 0,261,712; DE 195,04,224; WO 95/22586 and WO 97/00600.

[0026] Another object of the present invention is an anisotropic polymer film with charge transport properties obtainable from a polymerisable liquid crystal mixture as defined above that is aligned in its liquid crystal phase into macroscopically ordered orientation and polymerised or crosslinked to fix the oriented state.

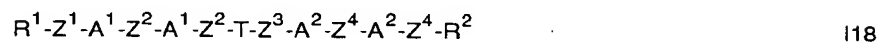
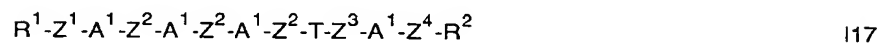
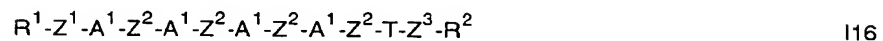
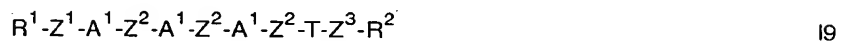
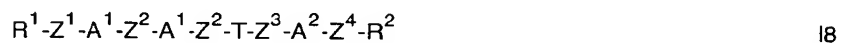
[0027] Particularly preferred are liquid crystal compounds of formula I, or liquid crystal mixtures comprising one or more compounds of formula I, that exhibit a nematic and/or smectic liquid crystal phase. For FET applications smectic materials are especially preferred. For OLED applications nematic or smectic materials are especially preferred.

[0028] Particularly preferred compounds of formula I are those wherein Z^1 , A^1 , Z^2 , T, Z^3 , A^2 and Z^4 form a conjugated system. Therein A^1 and A^2 are preferably arylene or heteroarylene and Z^1 , Z^2 , Z^3 and Z^4 are preferably a single bond or a conjugated link like for example $-C\equiv C-$ or $-CX^1=CX^2-$.

[0029] Further preferred compounds of formula I are those wherein

- m and o are 0,
- m and o are 1 or 2,
- T is dithienothiophene that may also be substituted with R^1 as defined above,
- n is 1 or 2 and Z^2 is a single bond, $-CX^1=CX^2-$ or $-C\equiv C-$.

[0030] Particularly preferred compounds of formula I are those of the following formulae



$R^1-Z^1-A^1-Z^2-A^1-Z^2-T-Z^3-T-Z^3-R^2$ I19

5 $R^1-Z^1-A^1-Z^2-A^1-Z^2-T-Z^2-A^1-Z^2-T-Z^3-R^2$ I20

$R^1-Z^1-A^1-Z^2-T-Z^2-A^1-Z^2-A^1-Z^2-T-Z^3-R^2$ I21

10 $R^1-Z^1-A^1-Z^2-A^1-Z^2-T-Z^3-T-Z^3-A^2-Z^4-R^2$ I22

15 $R^1-Z^1-A^1-Z^2-T-Z^2-A^1-Z^2-T-Z^3-A^2-Z^4-R^2$ I23

$R^1-Z^1-T-Z^2-A^1-Z^2-A^1-Z^2-A^1-Z^2-T-Z^3-R^2$ I24

20 $R^1-Z^1-A^1-Z^2-A^1-Z^2-T-Z^3-T-Z^3-T-Z^3-R^2$ I25

25 $R^1-Z^1-A^1-Z^2-T-Z^2-A^1-Z^2-T-Z^3-T-Z^3-R^2$ I26

$R^1-Z^1-A^1-Z^2-T-Z^2-T-Z^2-A^1-Z^2-T-Z^3-R^2$ I27

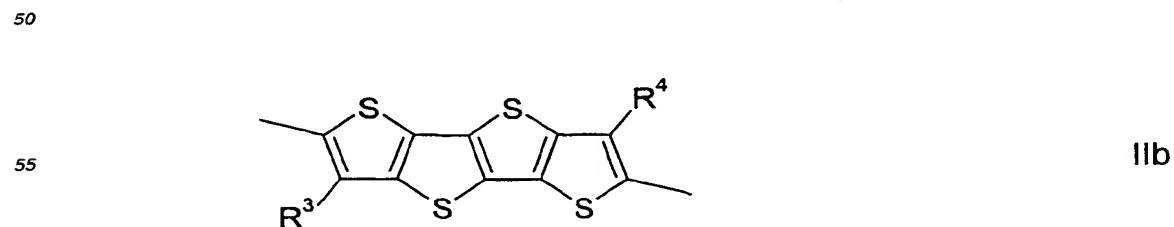
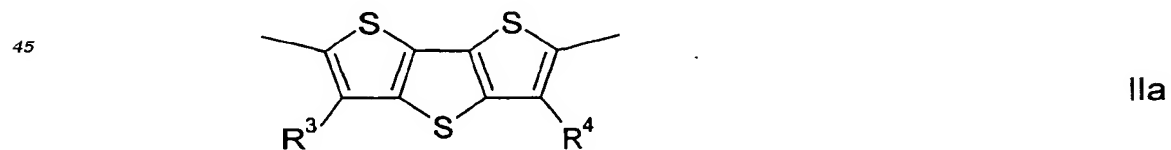
30 $R^1-Z^1-A^1-Z^2-T-Z^3-T-Z^3-T-Z^3-A^2-Z^4-R^2$ I28

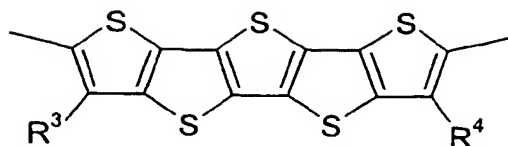
$R^1-Z^1-T-Z^2-A^1-Z^2-T-Z^2-A^1-Z^2-T-Z^3-R^2$ I29

35 $R^1-Z^1-T-Z^2-A^1-Z^2-A^1-Z^2-T^1-Z^3-T-Z^3-R^2$ I30

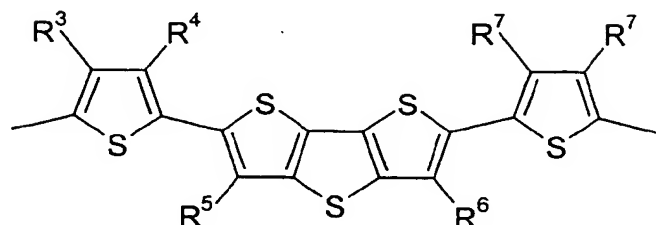
40 wherein R^1 , R^2 , Z^1 , Z^2 , Z^3 , Z^4 , A^1 , A^2 and T have, in case of multiple occurrence independently of each other, one of the meanings of formula I.

[0031] T is preferably selected from the following subformulae





IIc



IIId

wherein R^3 , R^4 , R^5 , R^6 , R^7 and R^8 have independently of each other one of the meanings of R^1 in formula I.

[0032] A^1 and A^2 are preferably selected from 1,4-phenylene, 1,4-cyclohexa-1,3-diene, 1,4-cyclohexenyne and in which, in addition, one or more CH groups may be replaced by N one or two non-adjacent CH_2 groups may be replaced by O and/or S, thiophene-2,5-diyl, thienothiophene-2,5-diyl, dithienothiophene-2,6-diyl, naphthalene-2,6-diyl, it being possible for all of these groups to be unsubstituted, mono- or polysubstituted by L, with L being halogen, CN, SCN, NO_2 , SF_5 or an alkyl, alkoxy, alkylcarbonyl, alkoxycarbonyl, silane or siloxy group with 1 to 12 C atoms, wherein one or more H atoms may be substituted with F or Cl.

[0033] A^1 and A^2 are particularly preferably 1,4-phenylene that is substituted with 1, 2 or 3 groups L as defined above, or thiophene-2,5-diyl, thienothiophene-2,5-diyl or dithienothiophene-2,6-diyl all of which are optionally substituted with one or more groups L as defined above.

[0034] Z^{1-4} are preferably selected from -O-, -S-, -OCH₂-, -CH₂O-, -SCH₂-, -CH₂S-, -CF₂O-, -OCF₂-, -CF₂S-, -SCF₂-, -CH₂CH₂-, -CF₂CH₂-, -CH₂CF₂-, -CF₂CF₂-, -CH=N-, -N=CH-, -N=N-, -CH=CR⁰-, -CX¹=CX²-, -C≡C- and a single bond, in particular from -CH=N-, -N=CH-, -N=N-, -CH=CR⁰-, -CX¹=CX²-, -C≡C- and a single bond.

[0035] Arylene and heteroarylene preferably denote a bivalent mono-, bi- or tricyclic aromatic or heteroaromatic group with up to 15 C atoms that may also comprise condensed rings and is optionally substituted with one or more groups R^1 . Very preferred arylene and heteroarylene groups are those having one of the preferred meanings of A^1 as given above and below.

[0036] -CX¹=CX²- is preferably -CH=CH-, -CH=CF-, -CF=CH-, -CF=CF-, -CH=C(CN)- or -C(CN)=CH-.

[0037] R^1 and R^2 in formula I are preferably alkyl or alkoxy with 1 to 25 C atoms that is optionally fluorinated.

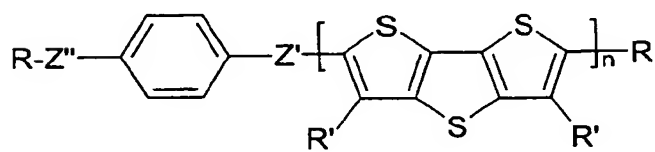
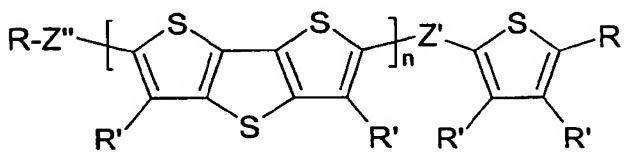
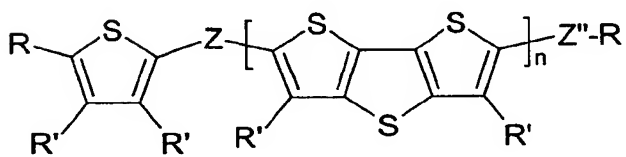
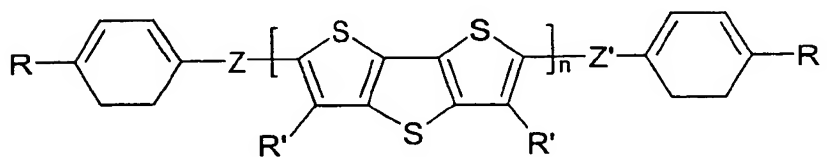
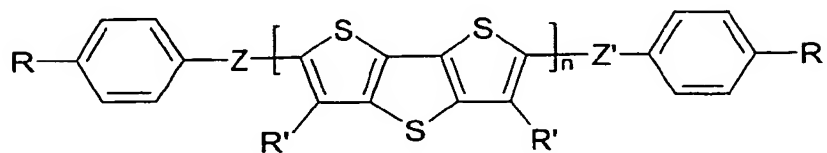
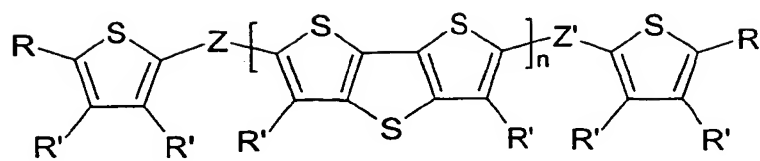
[0038] In the formulae shown above, R^1 to R^8 are preferably selected from C₁-C₂₀-alkyl, C₁-C₂₀-fluoroalkyl, C₁-C₂₀-alkenyl, C₁-C₂₀-alkynyl, C₁-C₂₀-alkoxy, C₁-C₂₀-thioether, C₁-C₂₀-silyl, C₁-C₂₀-ester and C₁-C₂₀-amino.

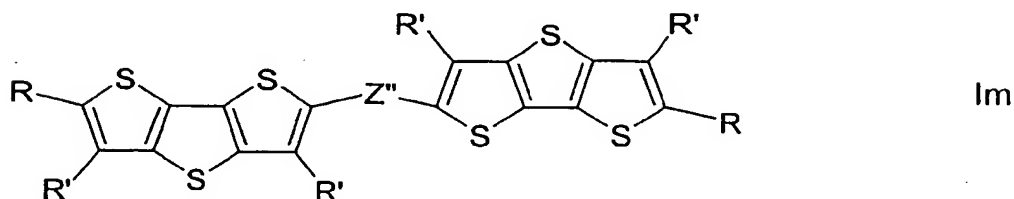
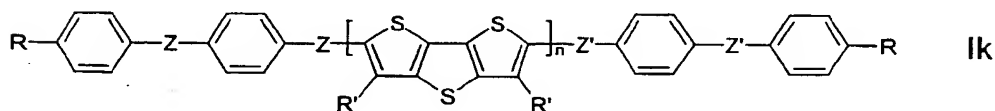
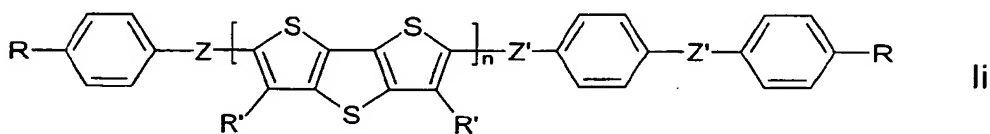
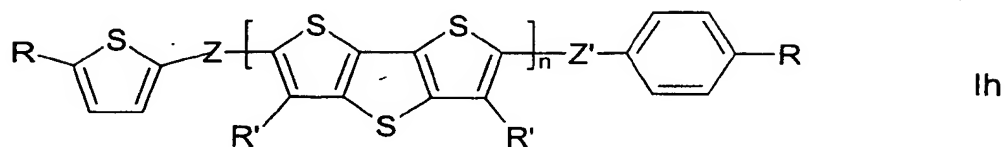
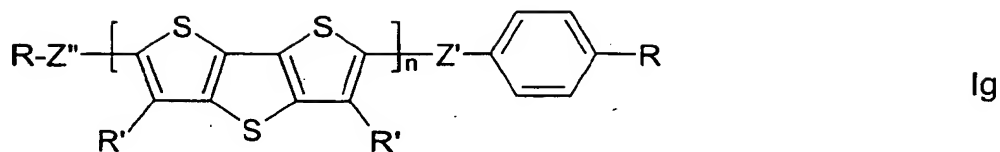
[0039] If one of R or R^1 to R^8 is an alkyl or alkoxy radical, i.e. where the terminal CH_2 group is replaced by -O-, this may be straight-chain or branched. It is preferably straight-chain, has 2, 3, 4, 5, 6, 7 or 8 carbon atoms and accordingly is preferably ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, ethoxy, propoxy, butoxy, pentoxy, hexoxy, heptoxy, or octoxy, furthermore methyl, nonyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, nonoxy, decoxy, undecoxy, dodecoxy, tridecoxy or tetradecoxy, for example.

[0040] Oxaalkyl, i.e. where one CH_2 group is replaced by -O-, is preferably straight-chain 2-oxapropyl (=methoxymethyl), 2- (=ethoxymethyl) or 3-oxabutyl (=2-methoxyethyl), 2-, 3-, or 4-oxapentyl, 2-, 3-, 4-, or 5-oxahexyl, 2-, 3-, 4-, 5-, or 6-oxaheptyl, 2-, 3-, 4-, 5-, 6- or 7-oxaoctyl, 2-, 3-, 4-, 5-, 6-, 7- or 8-oxanonyl or 2-, 3-, 4-, 5-, 6-, 7-, 8- or 9-oxadecyl, for example.

[0041] Halogen is preferably F or Cl.

[0042] Particularly preferred are the following compounds





wherein n has the meaning of formula I,

Z and Z' have independently of each other one of the meanings of Z¹ in formula II, and are preferably -CH=CH-, -CH=CF-, -CF=CH-, CH=CCl-, -CCl=CH-, -CF=CF-, -CCl=CCl-, -C≡C- or a single bond,

Z'' has one of the meanings of Z¹ in formula II, and is preferably -CH=CH-, -CH=CF-, -CF=CH-, CH=CCl-, -CCl=CH-, -CF=CF-, -CCl=CCl- or -C≡C-,

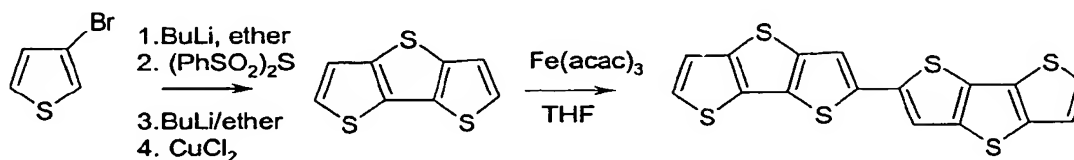
R and R' have independently of each other one of the meanings of R¹ given above, and are preferably halogen or an optionally fluorinated alkyl group with 1 to 15 C atoms.

[0043] In case one of the groups A¹⁻², Z¹⁻⁴, R¹⁻⁸, R, R', R⁰, R⁰⁰, Z', Z'' and T¹ appears more than once in a formula as shown above and below, the multiple groups may be identical or different, unless explicitly stated otherwise.

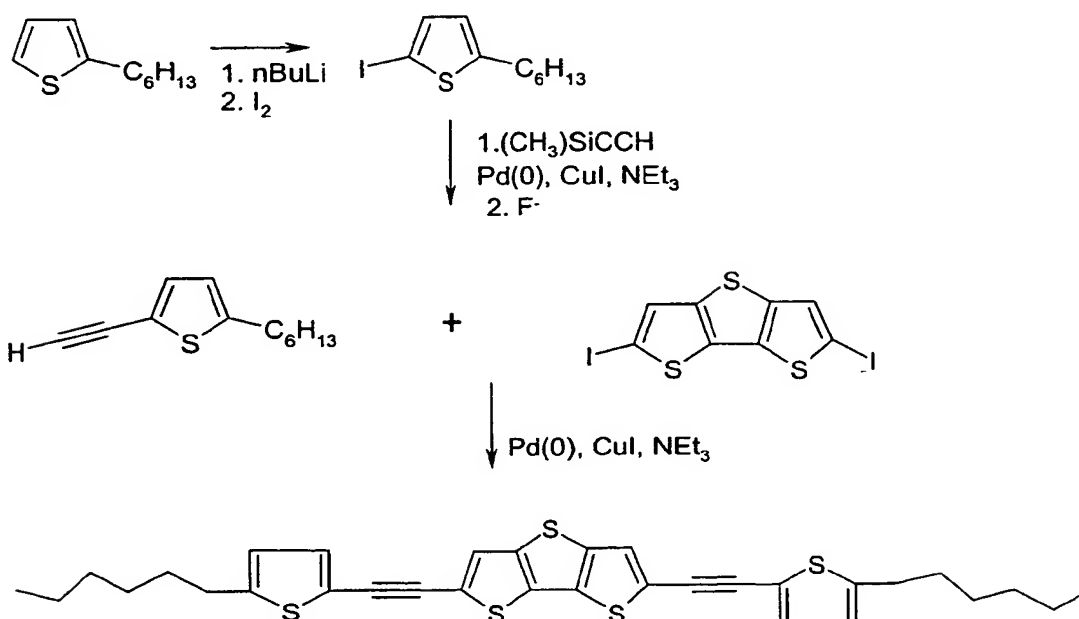
[0044] The compounds of formula I can be synthesized according to or in analogy to methods that are known to the skilled in the art and are described for example in F. de Jong and M. J. Janssen, J. Org. Chem., 1971, 36, 12, 1645;

S. Inaoka and D. M. Collard, J. Mater. Chem., 1999, 9, 1719 or WO 99/12989. Furthermore, they can be prepared according to or in analogy to the following reaction schemes. DTT and BTT can be prepared according to scheme 1.

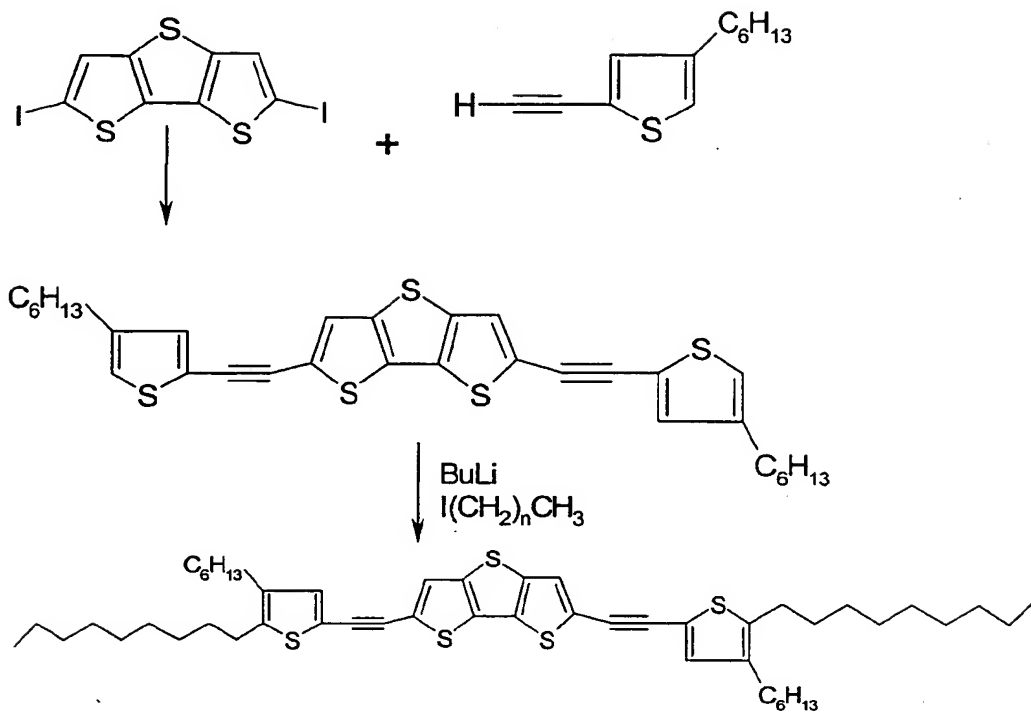
Scheme 1

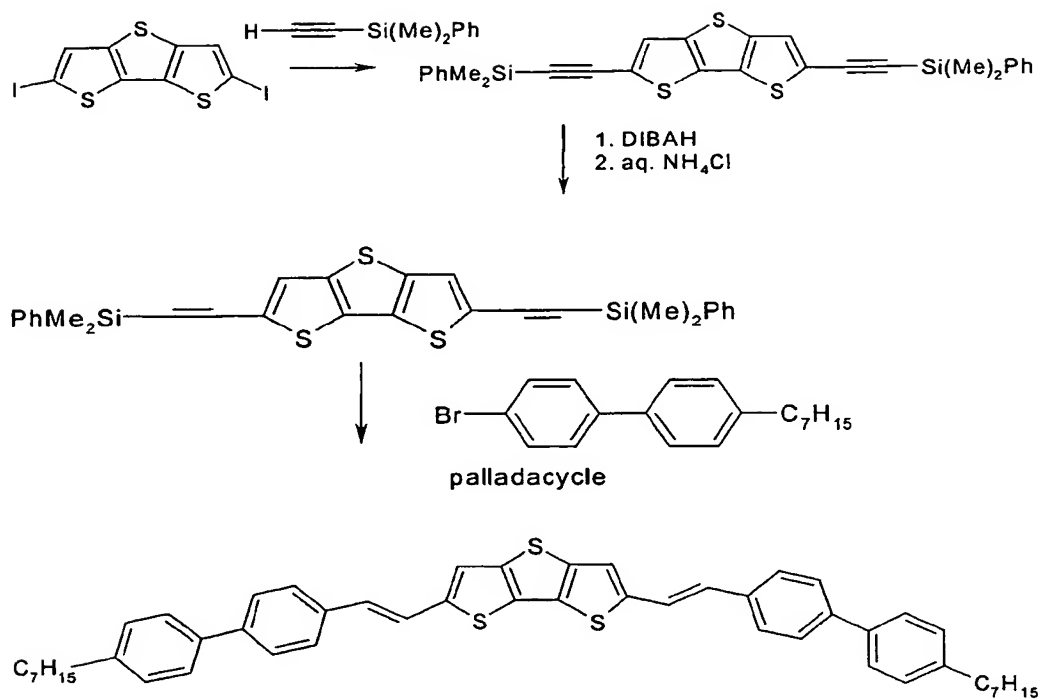


Scheme 2



Scheme 3



Scheme 4

10



30

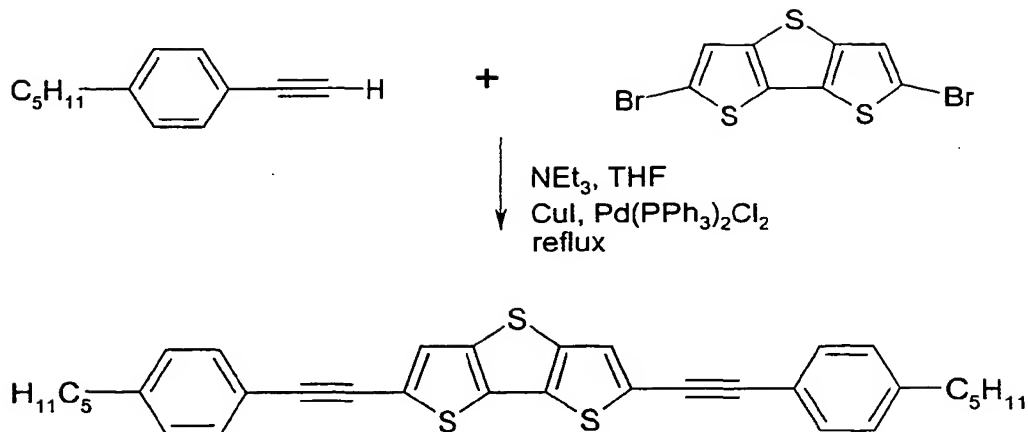


40

50

55

Scheme 7



[0045] A further aspect of the invention relates to both the oxidised and reduced form of the compounds and materials according to this invention. Either loss or gain of electrons results in formation of a highly delocalised ionic form, which is of high conductivity. This can occur on exposure to common dopants. Suitable dopants and methods of doping are known to those skilled in the art, e.g. from EP 0 528 662, US 5,198,153 or WO 96/21659.

[0046] The doping process typically implies treatment of the semiconductor material with an oxidating or reducing agent in a redox reaction to form delocalised ionic centres in the material, with the corresponding counterions derived from the applied dopants. Suitable doping methods comprise for example exposure to a doping vapor in the atmospheric pressure or at a reduced pressure, electrochemical doping in a solution containing a dopant, bringing a dopant into contact with the semiconductor material to be thermally diffused, and ion-implantation of the dopant into the semiconductor material.

[0047] When electrons are used as carriers, suitable dopants are for example halogens (e.g. I₂, Cl₂, Br₂, ICl, ICl₃, IBr and IF), Lewis acids (e.g. PF₅, AsF₅, SbF₅, BF₃, BCl₃, SbCl₅, BBr₃ and SO₃), protonic acids, organic acids, or amino acids (e.g. HF, HCl, HNO₃, H₂SO₄, HClO₄, FSO₃H and ClSO₃H), transition metal compounds (e.g. FeCl₃, FeOCl, Fe(CIO₄)₃, Fe(4-CH₃C₆H₄SO₃)₃, TiCl₄, ZrCl₄, HfCl₄, NbF₅, NbCl₅, TaCl₅, MoF₅, MoCl₅, WF₅, WCl₆, UF₆ and LnCl₃ (wherein Ln is a lanthanoid), anions (e.g. Cl⁻, Br⁻, I⁻, I₃⁻, HSO₄⁻, SO₄²⁻, NO₃⁻, ClO₄⁻, BF₄⁻, PF₆⁻, AsF₆⁻, SbF₆⁻, FeCl₄⁻, Fe(CN)₆³⁻, and anions of various sulfonic acids, such as aryl-SO₃⁻). When holes are used as carriers, examples of dopants are cations (e.g. H⁺, Li⁺, Na⁺, K⁺, Rb⁺ and Cs⁺), alkali metals (e.g., Li, Na, K, Rb, and Cs), alkaline-earth metals (e.g., Ca, Sr, and Br), O₂, XeOF₄, (NO₂⁺) (SbF₆⁻), (NO₂⁺) (SbCl₆⁻), (NO₂⁺) (BF₄⁻), AgClO₄, H₂IrCl₆, La (NO₃)₃·6H₂O, FSO₂OOSO₂F, Eu, acetylcholine, R₄N⁺ (R is an alkyl group), R₄P⁺ (R is an alkyl group), R₆As⁺ (R is an alkyl group), and R₃S⁺ (R is an alkyl group).

[0048] The conducting form of the compounds and materials of the present invention can be used as an organic "metal" in applications, for example, but not limited to, charge injection layers and ITO planarising layers in organic light emitting diode applications, films for flat panel displays and touch screens, antistatic films, printed conductive substrates, patterns of tracts in electronic applications such as printed circuit boards and condensers.

[0049] The mesogenic or liquid crystal compounds of formula I and the liquid crystal mixtures comprising compounds of formula I can be aligned in their liquid crystal state into homeotropic orientation, where the conjugated pi-electron systems are orthogonal to the direction of charge transport. This ensures that the intermolecular distances are minimised and hence then energy required to transport charge between molecules is minimised.

[0050] Alignment of the liquid crystal material can be achieved for example by treatment of the substrate onto which the material is coated, by shearing the material during or after coating, by application of a magnetic or electric field to the coated material, or by the addition of surface-active compounds to the liquid crystal material. Reviews of alignment techniques are given for example by I. Sage in "Thermotropic Liquid Crystals", edited by G. W. Gray, John Wiley & Sons, 1987, pages 75-77, and by T. Uchida and H. Seki in "Liquid Crystals - Applications and Uses Vol. 3", edited by B. Bahadur, World Scientific Publishing, Singapore 1992, pages 1-63. A review of alignment materials and techniques is given by J. Cognard, Mol. Cryst. Liq. Cryst. 78, Supplement 1 (1981), pages 1-77.

[0051] In case of polymerisable materials, polymerisation can be achieved by exposure to heat or actinic radiation. Actinic radiation means irradiation with light, like UV light, IR light or visible light, irradiation with X-rays or gamma rays

or irradiation with high energy particles, such as ions or electrons. Preferably polymerisation is carried out by UV irradiation at a non-absorbing wavelength. As a source for actinic radiation for example a single UV lamp or a set of UV lamps can be used. When using a high lamp power the curing time can be reduced. Another possible source for actinic radiation is a laser, like e.g. a UV laser, an IR laser or a visible laser.

[0052] Polymerisation is preferably carried out in the presence of an initiator absorbing at the wavelength of the actinic radiation. For example, when polymerising by means of UV light, a photoinitiator can be used that decomposes under UV irradiation to produce free radicals or ions that start the polymerisation reaction. When curing polymerisable materials with acrylate or methacrylate groups, preferably a radical photoinitiator is used, when curing polymerisable materials with vinyl, epoxide and oxetane groups, preferably a cationic photoinitiator is used. It is also possible to use a polymerisation initiator that decomposes when heated to produce free radicals or ions that start the polymerisation. As a photoinitiator for radical polymerisation for example the commercially available Irgacure 651, Irgacure 184, Darocure 1173 or Darocure 4205 (all from Ciba Geigy AG) can be used, whereas in case of cationic photopolymerisation the commercially available UVI 6974 (Union Carbide) can be used.

[0053] The polymerisable material can additionally comprise one or more other suitable components such as, for example, catalysts, sensitizers, stabilizers, inhibitors, chain-transfer agents, co-reacting monomers, surface-active compounds, lubricating agents, wetting agents, dispersing agents, hydrophobing agents, adhesive agents, flow improvers, defoaming agents, deaerators, diluents, reactive diluents, auxiliaries, colourants, dyes or pigments.

[0054] The compounds of formula I and the mixtures obtained thereof are useful as optical, electronic and semiconductor materials, in particular as charge transport materials in field effect transistors (FETs) e.g. as components of integrated circuitry, ID tags or TFT applications. Alternatively, they may be used in organic light emitting diodes (OLEDs) in electroluminescent display applications or as backlight of e.g. liquid crystal displays, as photovoltaics or sensor materials, and for other semiconductor applications, as electrode materials in batteries, as photoconductors and for electrophotographic applications like electrophotographic recording.

[0055] FETs comprising compounds of formula I or mixtures or polymers comprising them are suitable for example as ID tags, containing specific information in clothing, food containers and other consumer products. In security applications they are suitable for use in field effect transistors for ID tags or security markings to authenticate and prevent counterfeiting of documents of value like banknotes, credit cards or ID cards, national ID documents, licenses or any product with money value, like stamps, tickets, shares, cheques etc..

[0056] The materials of the present invention are useful as optical, electronic and semiconductor materials, in particular as charge transport materials in field effect transistors (FETs), as photovoltaics or sensor materials, for electrophotographic recording, and for other semiconductor applications. Such FETs, where an organic semiconductive material is arranged as a film between a gate-dielectric and a drain and a source electrode, are generally known e.g. from US 5,892,244, WO 00/79617, US 5,998,804, and from the references cited in the background and prior art chapter. Due to the advantages, like low cost production using the solubility properties of the compounds according to the invention and thus the processibility of large surfaces, preferred applications of these FETs are such as integrated circuitry, TFT-displays and security applications.

[0057] Alternatively, the materials according to the invention may be used in organic light emitting devices or diodes (OLEDs), e.g. in display applications or as backlight of e.g. liquid crystal displays. Common OLEDs are realized using multilayer structures. An emission layer is generally sandwiched between one or more electron-transport and/or hole-transport layers. By applying an electric voltage electrons and holes as charge carriers move towards the emission layer where their recombination leads to the excitation and hence luminescence of the lumophor units contained in the emission layer. The inventive compounds, materials and films may be employed in one or more of the charge transport layers and/ or in the emission layer, corresponding to their electrical and/ or optical properties. Furthermore their use within the emission layer is especially advantageous, if the compounds, materials and films according to the invention show electroluminescent properties themselves or comprise electroluminescent groups or compounds. The selection, characterization as well as the processing of suitable monomeric, oligomeric and polymeric compounds or materials for the use in OLEDs is generally known by a person skilled in the art, see e. g. Meerholz, *Synthetic Materials*, 111-112, 2000, 31-34, Alcalá, *J. Appl. Phys.*, 88, 2000, 7124-7128 and the literature cited therein.

[0058] According to another use, the inventive compounds, materials or films, especially those which show photoluminescent properties, may be employed as materials of light sources, e.g. of display devices such as described in EP 0 889 350 A1 or by C. Weder et al., *Science*, 279, 1998, 835-837.

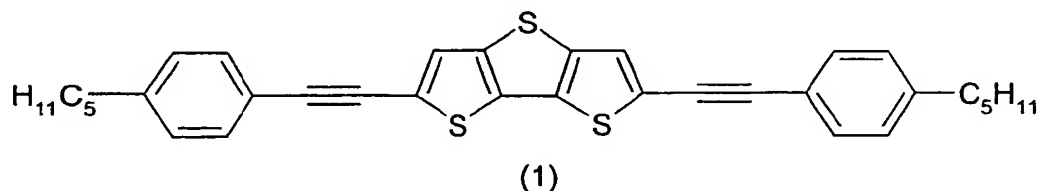
[0059] Furthermore, the compounds of the present invention are useful as high birefringence compounds added to liquid crystalline compositions in order to increase birefringence. For this purpose, they do not need to have a mesophase themselves, but a similar shape to conventional liquid crystals in order to dissolve and not to detract from the liquid crystal properties of the composition.

[0060] In the foregoing and in the following examples, unless otherwise indicated, all temperatures are set forth uncorrected in degrees Celsius and all parts and percentages are by weight. The following abbreviations are used to illustrate the liquid crystalline phase behaviour of the compounds: K = crystalline; N = nematic; S = smectic; Ch =

cholesteric; I = isotropic. The numbers between the symbols indicate the phase transition temperatures in °C.

Example 1

[0061] 2,6-Bis-(4-pentyl-phenylethynyl)-dithieno[3,2-*b*:2',3'-*d*]thiophene (1) was prepared according to scheme 7 and as described below



[0062] 2,6-Dibromo-dithieno[3,2-*b*:2',3'-*d*]thiophene was prepared according to a procedure described in G. F. Pedulli, M. Tiecco, M. Guera, G. Martelli and P. Zanirato, *J. C. S. Perkin II*, 1978, 212. 2,6-Dibromo-dithieno[3,2-*b*:2',3'-*d*]thiophene (0.5 g, 1.4 mmol), triethylamine (15 ml), and a catalytic amount of palladium bis(triphenylphosphine) dichloride and copper iodide were stirred under nitrogen in tetrahydrofuran. The solution was warmed to 60°C and 4-pentyl-phenylacetylene (1.1 g, 6.4 mmol) dissolved in tetrahydrofuran (30 ml) was added dropwise over a period of 2 hours. The solution was heated under reflux overnight. The brown solution was poured in to dichloromethane, washed with water, the chlorinated phase was removed, dried over sodium sulphate and evaporated to dryness. The residue was purified by flash column chromatography using petroleum (80-100) followed by dichloromethane as eluant.

[0063] Evaporation of the appropriate fractions yielded (1) as a bright yellow solid (320 mg). ¹H NMR and ¹³C NMR showed expected signals.

[0064] The following transitions and phases were observed by optical microscopy:

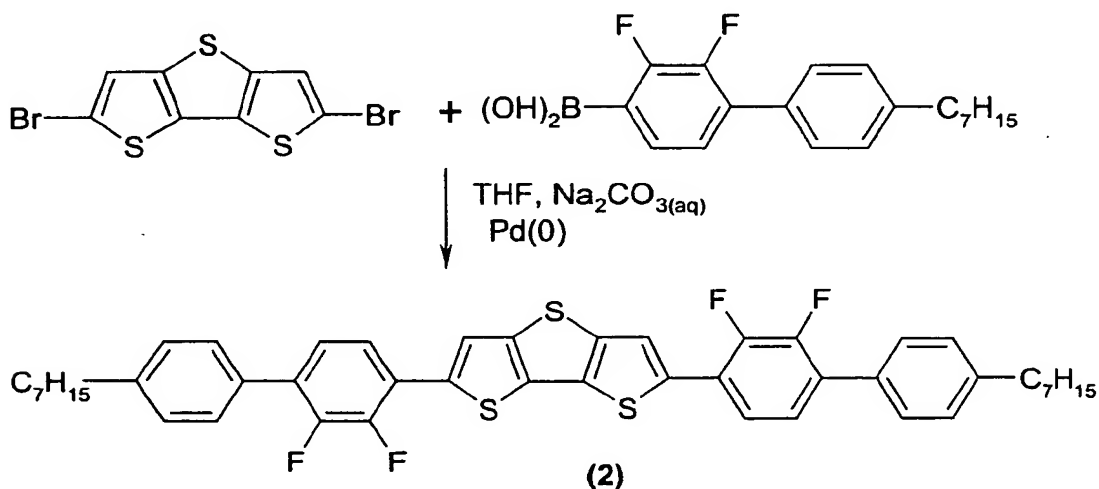
K-134-N-235-I (first heat)

I-234-N-110-K (first cool)

K-134-N-234-I (second heat)

Example 2

[0065] Compound (2) was prepared as follows:



[0066] 2,6-Dibromo-dithieno[3,2-*b*;2',3'-*d*]thiophene (1.0 g, 2.8 mmol) and 4'-heptylphenyl-2,3-difluorophenyl benzoic acid (2.1 g), sodium carbonate (1.07 g, 10.1 mmol in 20 ml water) and a catalytic amount of palladium bis(triphenylphosphine) dichloride were stirred under reflux in tetrahydrofuran. After 16 h, the brown solution was poured in to dichloromethane, washed with water, the chlorinated phase was removed, dried over sodium sulphate and evaporated to dryness. The residue was purified by flash column chromatography using petroleum (80-100) followed by dichloromethane as eluant. Evaporation of the appropriate fractions yielded (2) as a bright yellow solid (430 mg). ¹H and ¹³C NMR showed expected signals.

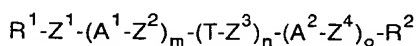
[0067] The following transitions and phases were observed by optical microscopy:

K-123-N-262-I (first heat)

I-256-N-112-K (first cool)

Claims

1. Thienothiophenes of formula I



wherein

R¹ and R² are independently of each other H, halogen, CN, NO₂, straight chain, branched or cyclic alkyl with 1 to 20 C-atoms, which may be unsubstituted, mono- or poly-substituted by F, Cl, Br, I or CN, it being also possible for one or more non-adjacent CH₂ groups to be replaced, in each case independently from one another, by -O-, -S-, -NH-, -NR⁰-, -SiR⁰R⁰⁰-, -CO-, -COO-, -OCO-, -OCO-O-, -S-CO-, -CO-S-, -CH=CH- or -C≡C- in such a manner that O and/or S atoms are not linked directly to one another,

R⁰ and R⁰⁰ are independently of each other H or alkyl with 1 to 12 C-atoms,

A¹ and A² are independently of each other an alicyclic or aromatic group that may also comprise one or more hetero atoms and one or more fused rings, and A¹ may also denote T,

Z¹ to Z⁴ are independently of each other -O-, -S-, -CO-, -COO-, -OCO-, -O-COO-, -CO-NR⁰-, -NR⁰-CO-, -OCH₂-, -CH₂O-, -SCH₂-, -CH₂S-, -CF₂O-, -OCF₂-, -CF₂S-, -SCF₂-, -CH₂CH₂-, -CF₂CH₂-, -CH₂CF₂-, -CF₂CF₂-, -CH=N-, -N=CH-, -N=N-, -CH=CR⁰-, -CX¹=CX²-, -C≡C-, -CH=CH-COO-, -OCO-CH=CH- or a single bond,

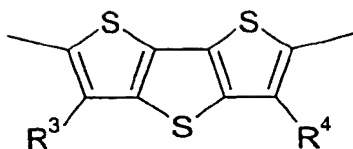
X¹ and X² are independently of each other H, F, Cl or CN,

T is a group consisting of 3, 4, 5 or 6 fused thiophene rings which may also be mono- or polysubstituted by R¹,

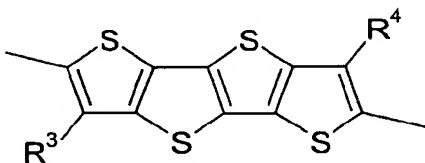
m and o are independently of each other 0, 1, 2 or 3, and

n is 1, 2 or 3.

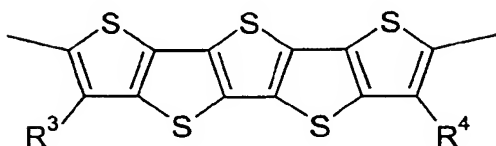
2. Thienothiophenes according to claim 1, wherein T is selected from the following subformulae



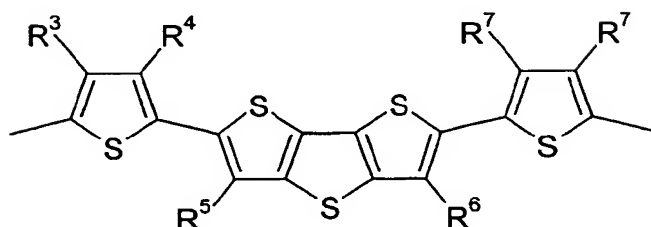
IIa



IIb



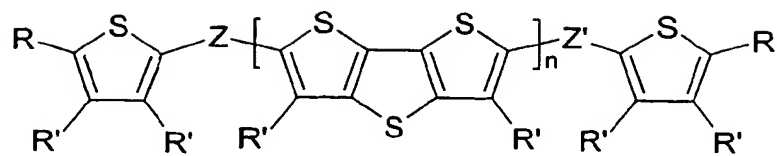
IIc



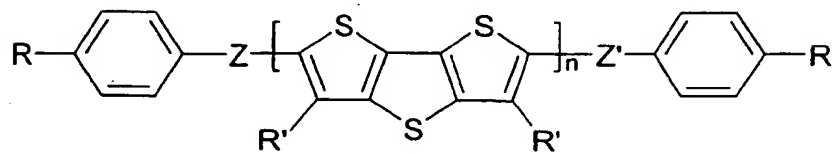
IIId

wherein R^3 , R^4 , R^5 , R^6 , R^7 and R^8 have independently of each other one of the meanings of R^1 in formula I.

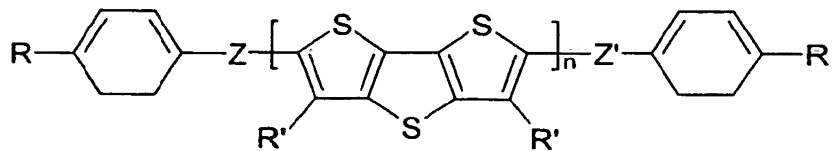
3. Thienothiophenes according to claim 1 or 2, wherein A^1 and A^2 are selected from 1,4-phenylene, 1,4-cyclohexa-1,3-diene, 1,4-cyclohexenylene, in which, in addition, one or more CH groups may be replaced by N one or two non-adjacent CH_2 groups may be replaced by O and/or S, thiophene-2,5-diyl, thienothieophene-2,5-diyl, dithienothiophene-2,6-diyl, furan-2,5-diyl, 1,4-bicyclo-(2,2,2)-octylene, naphthalene-2,6-diyl, and indane-2,5-diyl, it being possible for all of these groups to be unsubstituted, mono- or polysubstituted by L, with L being halogen, CN, SCN, NO_2 , SF_5 or an alkyl, alkoxy, alkylcarbonyl or alkoxy carbonyl group with 1 to 4 C atoms, wherein one or more H atoms may be substituted with F or Cl.
4. Thienothiophenes according to at least one of claims 1 to 3, wherein R^1 to R^8 are selected from H, F, Cl, CN, NO_2 , straight chain, branched or cyclic alkyl with 1 to 20 C-atoms, which may be unsubstituted, mono- or poly-substituted by F, Cl, Br, I or CN, it being also possible for one or more non-adjacent CH_2 groups to be replaced, in each case independently from one another, by -O-, -S-, -NH-, - NR^0 -, - SiR^0R^0 -, -CO-, -COO-, -OCO-, -OCO-O-, -S-CO-, -CO-S-, -CH=CH- or -C≡C- in such a manner that O and/or S atoms are not linked directly to one another.
5. Thienothiophenes according to at least one of claims 1 to 4, wherein Z^1 , Z^2 , Z^3 and Z^4 are selected from -CH=CH-, -CH=CF-, -CF=CH-, -CF=CF-, -CH=C(CN)- or -C(CN)=CH-, -C≡C- and a single bond.
6. Thienothiophenes according to at least one of claims 1 to 5, selected from the following formulae



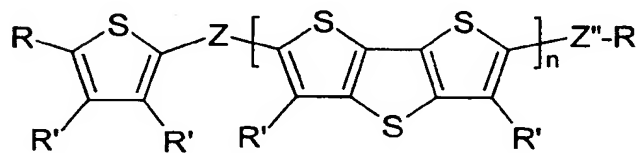
la



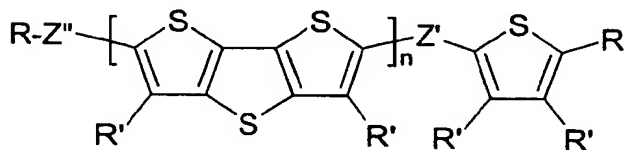
lb



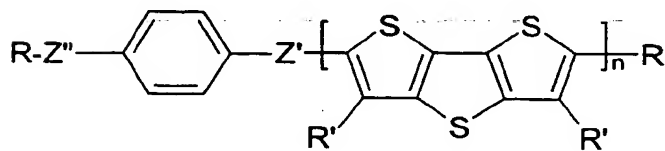
lc



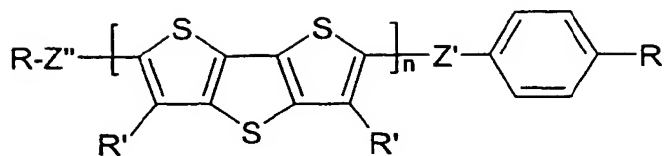
ld



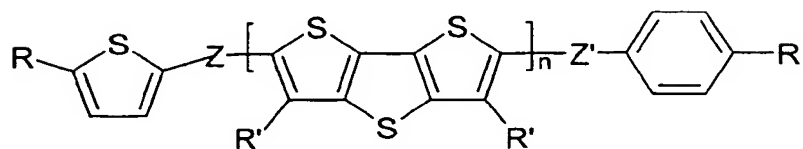
le



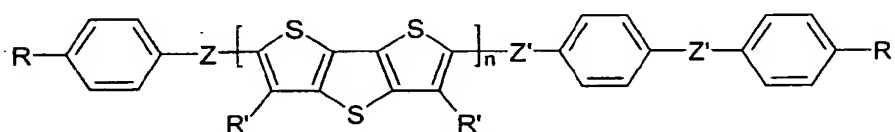
lf



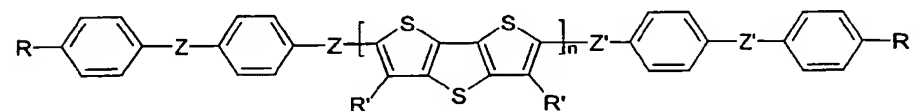
lg



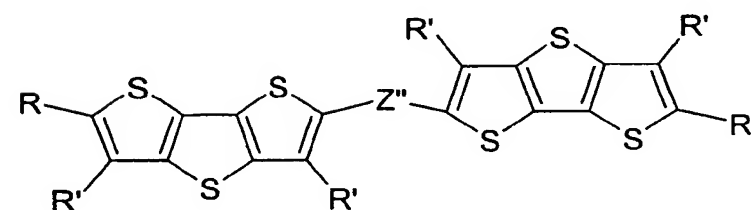
lh



li



lk



lm

wherein n has the meaning of formula I,

Z and Z' have independently of each other one of the meanings of Z¹ in formula II, and are preferably -CH=CH-, -CH=CF-, -CF=CH-, CH=CCl-, -CCl=CH-, -CF=CF-, -CCl=CCl-, -C≡C- or a single bond,

Z'' has one of the meanings of Z¹ in formula II, and is preferably -CH=CH-, -CH=CF-, -CF=CH-, CH=CCl-, -CCl=CH-, -CF=CF-, -CCl=CCl- or -C≡C-,

R and R' have independently of each other one of the meanings of R¹ given above, and are preferably halogen or an optionally fluorinated alkyl group with 1 to 15 C atoms.

7. Thienothiophenes according to at least one of claims 1 to 6 that are mesogenic or liquid crystalline.

8. Liquid crystal mixture comprising one or more thienothiophenes according to at least one of claims 1 to 7 and

optionally one or more further compounds, wherein at least one of said thienothiophenes and further compounds is mesogenic or liquid crystalline.

- 5
9. Use of thienothiophenes of claims 1 to 7 or a mixture of claim 8 as semiconductors or charge transport materials in optical, electrooptical or electronic devices, like for example components of integrated circuitry, field effect transistors (FET) for example as thin film transistors in flat panel display applications or for Radio Frequency Identification (RFID) tags, and in semiconducting components for organic light emitting diode (OLED) applications, electroluminescent display devices, backlights, photovoltaic or sensor devices, as electrode materials in batteries, as photoconductors and for electrophotographic applications.
- 10
10. Field effect transistor, OLED, electroluminescent device, RFID tag, backlight, photovoltaic or sensor device, photoconductor and electrophotographic recording device comprising a thienothiophene of claims 1 to 7 or a mixture of claim 8.
- 15
11. Security marking or device comprising a thienothiophene of claims 1 to 7, a mixture of claim 8 or a FET or RFID according to claim 10.
- 20
12. Compounds or a mixture according to at least one of claims 1 to 8, which are oxidatively or reductively doped to form conducting ionic species.
- 25
13. Charge injection layer, planarising layer, antistatic film or conducting substrate or pattern for electronic applications or flat panel displays, comprising compounds or a mixture according to claim 12.
- 30
- 35
- 40
- 45
- 50
- 55

THIS PAGE BLANK (USPTO)

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 275 651 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
14.05.2003 Bulletin 2003/20

(51) Int Cl.7: **C07D 495/14**, C07D 519/00,
C09K 19/34, H01B 1/12

(43) Date of publication A2:
15.01.2003 Bulletin 2003/03

(21) Application number: **02013425.0**

(22) Date of filing: **13.06.2002**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **09.07.2001 EP 01115742**

(71) Applicant: **MERCK PATENT GmbH**
64293 Darmstadt (DE)

(72) Inventors:
• **Farrand, Louise**
Blandford Forum, Dorset DT11 9ED (GB)

- **Thompson, Marcus**
Fordingbridge, Hampshire SP6 1RR (GB)
- **Giles, Mark**
Southampton SO15 2LE (GB)
- **Heeney, Martin**
Southampton SO14 6TQ (GB)
- **Tierney, Steven**
Southampton SO15 7QW (GB)
- **Shkunov, Maxim**
Southampton SO16 6SX (GB)
- **Sparrowe, David**
Bournemouth, Dorset BH6 5EJ (GB)
- **McCulloch, Iain**
Kings Somborne, Hampshire SO20 6PE (GB)

(54) **Thienothiophene derivatives**

(57) The invention relates to new thienothiophene derivatives, their use as semiconductors or charge transport materials, in optical, electrooptical or electronic devices like for example organic field effect transistors (FET or OFET) for thin film transistor liquid crystal displays and integrated circuit devices such as RFID tags,

electroluminescent devices in flat panel displays, and in photovoltaic and sensor devices, and to a field effect transistor, light emitting device or ID tag comprising the thienothiophene derivatives.

EP 1 275 651 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 02 01 3425

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
D,A	WO 99 12989 A (CAMBRIDGE DISPLAY TECHNOLOGY LTD.) 18 March 1999 (1999-03-18) * the whole document *	1-13	C07D495/14 C07D519/00 C09K19/34 H01B1/12
A	KOSSMEHL G ET AL: "Über Polyarylenalkenylene und Polyheteroarylenalkenylene" MAKROMOLEKULARE CHEMIE, MACROMOLECULAR CHEMISTRY AND PHYSICS, HUTHIG UND WEPF VERLAG, BASEL, CH, vol. 183, no. 11, 1 November 1982 (1982-11-01), pages 2771-2786, XP002085568 ISSN: 0025-116X * the whole document *	1-13	
A	HARISTOY D ET AL: "STRUCTURE AND PHOTOCONDUCTIVE BEHAVIOUR OF A SANIDIC LIQUID CRYSTAL" LIQUID CRYSTALS, TAYLOR AND FRANCIS LTD, LONDON, GB, vol. 27, no. 3, March 2000 (2000-03), pages 321-328, XP000932241 ISSN: 0267-8292 * the whole document *	1-13	TECHNICAL FIELDS SEARCHED (Int.Cl.7) C07D C09K H01B
A	KIEBOOMS R ET AL: "SYNTHESIS, ELECTRICAL, AND OPTICAL PROPERTIES OF CONJUGATED POLYMERS" HANDBOOK OF ADVANCED ELECTRONIC AND PHOTONIC MATERIALS AND DEVICES, XX, XX, vol. 8, 2001, pages 1-102, XP001029240 * the whole document *	1-13	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 20 March 2003	Examiner Beslier, L
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

EPO FORM 1503 03 82 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 02 01 3425

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

20-03-2003

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9912989 A	18-03-1999	AU 8991698 A	29-03-1999
		EP 1017735 A1	12-07-2000
		WO 9912989 A1	18-03-1999
		JP 3313357 B2	12-08-2002
		JP 2001515933 T	25-09-2001
		US 6403809 B1	11-06-2002
<hr/>			

EPOFORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

THIS PAGE BLANK (USPTO)